

# Barriers to and opportunities for improved MASLD/MASH education: a qualitative discussion with medical training program leaders

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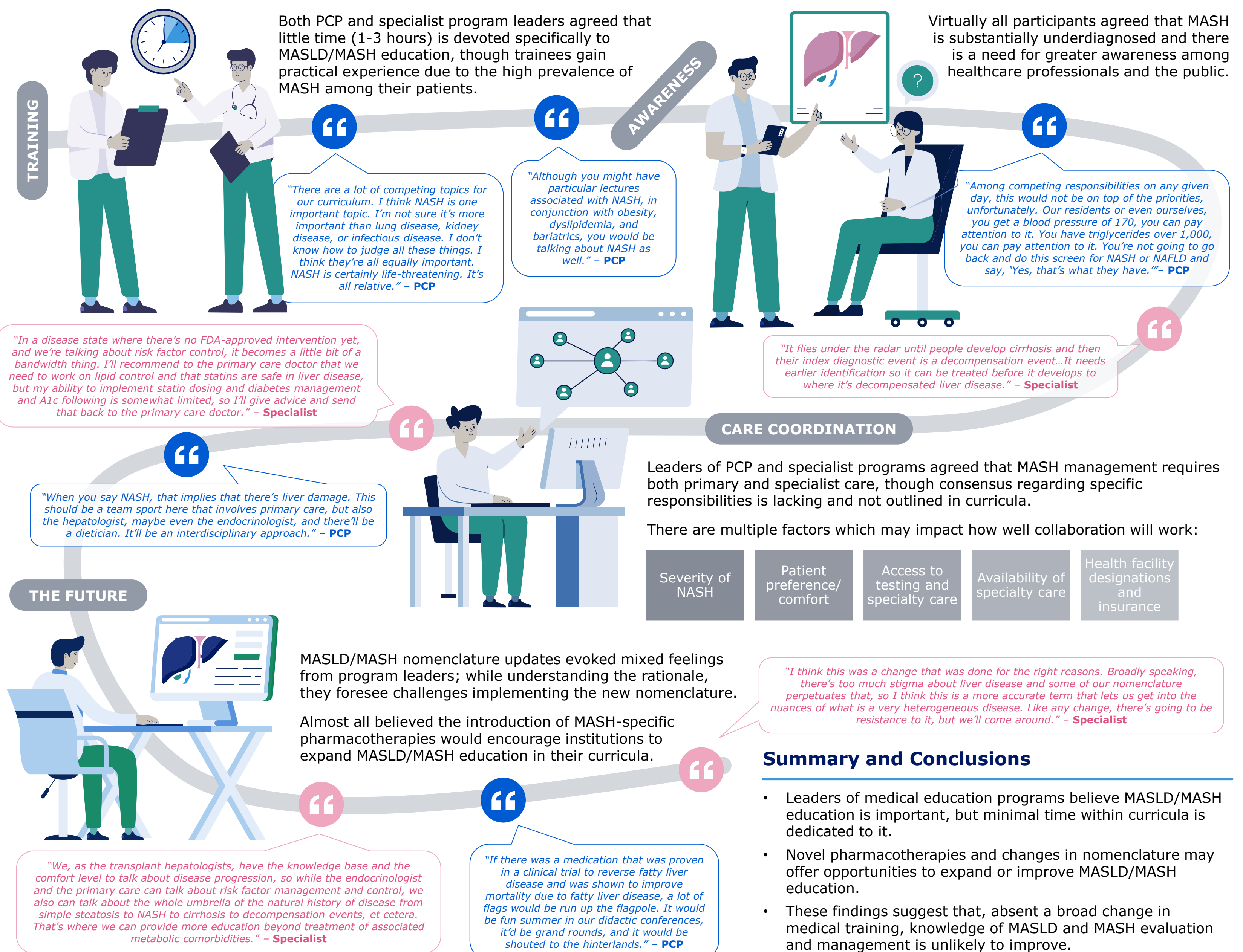
## Background and Aim

- Metabolic dysfunction-associated steatotic liver disease (MASLD, formerly NAFLD) and metabolic dysfunction-associated steatohepatitis (MASH, formerly NASH) are increasingly prevalent in the United States.
- However, MASLD and MASH remain substantially underdiagnosed, and many patients with MASH are not receiving care in accordance with the latest guidelines.
- We aimed to understand how MASLD/MASH-specific training is currently offered in medical training programs and to identify barriers to and opportunities for the improvement of MASLD/MASH education.

## Methods

- We conducted two qualitative 90-minute virtual focus groups with leaders of medical training programs on August 24 and 28, 2023.
- One group included leaders of primary care (nurse practitioner/physician assistant, internal medicine, family medicine) programs (PCP, n = 5).
- The second group included leaders of specialist (endocrinology, gastroenterology, hepatology) programs (n = 6).
- PCP program leaders reported 4 to 23 years in their current roles; specialist program leaders reported 3 to 16 years.
- Participants were recruited by email from a pool of participants who participated previously in a larger quantitative survey on MASH curricula.
- They were knowledgeable about their curricula (self-reported) and only one participant per institution was allowed.
- An institutional review board (IRB) exemption was issued.

## Results



## Summary and Conclusions

- Leaders of medical education programs believe MASLD/MASH education is important, but minimal time within curricula is dedicated to it.
- Novel pharmacotherapies and changes in nomenclature may offer opportunities to expand or improve MASLD/MASH education.
- These findings suggest that, absent a broad change in medical training, knowledge of MASLD and MASH evaluation and management is unlikely to improve.

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This study was sponsored by Novo Nordisk Inc. The authors acknowledge the medical writing assistance of John Newman, PhD, and data analysis of Andrea Stoltz, MS, of KJT Group, Inc. (Rochester, NY, USA).

Presented at NASH-TAG, Park City, UT (Jan. 4-6, 2024).

Disclosures:

AH, AA, and TF are employees and shareholders of Novo Nordisk Inc., which funded this research.

AMA serves on the advisory board for and receives speaking/teaching fees from Novo Nordisk. She receives research grants from Target Pharma and Pfizer.

MC has consulted for Novo Nordisk, 89Bio, Sagimet, Galecto, Cytodyn, Pfizer, Bristol Myers Squibb, Merck, Gilead, Akero, Theratechnologies, and Northsea and provided research support for Northsea, Pfizer, and Novo Nordisk.

SH has worked with AgomAB, Akero Therapeutics, Inc, Alentis Therapeutics AG, Aligos Therapeutics, Inc, Alimentiv, Inc, Altimune, Inc, Arrowhead Pharmaceuticals, Inc, Axcella Health, Inc, Blade Therapeutics, Bluejay Therapeutics, BMS, Boston Pharmaceuticals, Boxer Capital, BVF Partners, Can-Fite Biopharma Ltd, Chronwell, Inc, Cirus Therapeutics, Inc, Civi Biopharma, Inc, Cldf, CohBar, Inc, Conatus, Concept, Cymabay Therapeutics, Inc, Echosens North America, Inc, Enyo Pharma S.A., Fironostics, Inc, Foresite Labs, LLC, Fortress Biotech, Inc, Galectin Therapeutics, Inc, Galecto, Inc, Galmed Research & Dev, Ltd, Gelesis, Inc, Genentech (subsidiary of Roche), Genfit Corp, Gilead Sciences, Inc, Glaxo Smith Kline (GSK), GNS Healthcare, Inc, GRI Bio, Hepagene Therapeutics, Hepion Pharmaceuticals, Inc, Hepta Bio Inc, Hightide Therapeutics, Inc, HistoIndex Pte. Ltd., Humana, Immuron, Indalo Therapeutics, Inc, Inipharm, Inc, Innovate Biopharmaceuticals, Inc, Intercept Pharmaceuticals, Inc, Ionis, Kowa Research Institute, Inc, Madrigal Pharmaceuticals, Inc, Medpace, Inc, Merck Sharp & Dohme Corp, Metacrine, Inc, MGGM Therapeutics LLC, NeuroBo

Pharmaceuticals Inc, NGM Biopharmaceuticals, Inc, NorthSea Therapeutics B.V., Novartis Pharmaceuticals Corp, Novo Nordisk, Nutrasource, PathAI, Inc, Perspectum Diagnostics, Inc, Pfizer, Piper Sandler & Co, Poxel, Prometic (now Liminal), Ridgeline Therapeutics, Sagimet Biosciences, Second Genome, Inc, Silverback Therapeutics, Inc, Sonic Incytes Medical Corp, Terns, Inc, Viking Therapeutics, Inc, and Zafgen

DD has consulted for Novo Nordisk and been a speaker for Intercept.

MN serves on the advisory board for Altimune, BI, Cytodyn, 89Bio, GSK, Madrigal, Merck, Novo Nordisk, Perspectum, Siemens, Terns and Takeda. He is a principal investigator for a drug study with Allergan, Akero, BMS, Gilead, Galactin, Genfit, GSK, Conatus, Concept, Enanta, Madrigal, Novartis, Novo Nordisk, Shire, Takeda, Terns, Viking and Zyudus. He is a stockholder with Rivus Pharma, CIMa, Cytodyn, and ChronWell.